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L6
    ANSWER 1 OF 1 REGISTRY COPYRIGHT 2006 ACS on STN
RN
    511274-47-8 REGISTRY
CN
     D-Glucitol, 1,5-anhydro-1-C-phenyl-4,6-O-[(R)-phenylmethylene]-, (1S)-
     (9CI) (CA INDEX NAME)
FS
     STEREOSEARCH
MF
     C19 H20 O5
SR
                 CA, CAPLUS, TOXCENTER, USPAT2, USPATFULL
LC
     STN Files:
DT.CA Caplus document type: Journal; Patent
      Roles from patents: BIOL (Biological study); PREP (Preparation); RACT
       (Reactant or reagent); USES (Uses)
RL.NP Roles from non-patents: PREP (Preparation); RACT (Reactant or reagent)
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Absolute stereochemistry. Rotation (+).

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 4 REFERENCES IN FILE CA (1907 TO DATE)
- 4 REFERENCES IN FILE CAPLUS (1907 TO DATE)

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E3
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F.4
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E11
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E12
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The indicated field code is not available for EXPAND in this
file. To see a list of valid EXPAND field codes, enter HELP
SFIELDS at an arrow prompt (=>).
=> s e4-e6
'IN' IS NOT A VALID FIELD CODE
            47 ("SAS BENEDIKT"/IN OR "SAS BENEDIKT B 2300 TURNHOUT BE"/IN OR
               "SAS BENEDIKT J"/IN)
=> s l1 and DIARRHEA(s) VIRUSES
             5 L1 AND DIARRHEA(S) VIRUSES
=> d ibib 1-5
       ANSWER 1 OF 5
                         EPFULL
                                 COPYRIGHT 2006 EPO/FIZ KA on STN
                        2003:139988 EPFULL
ACCESSION NUMBER:
ENTRY DATE PUBLICATION: 20051102
UPDATE DATE PUBLICAT.: 20051102
DATA UPDATE DATE:
                        20051102
DATA UPDATE WEEK:
                     200544
TITLE (ENGLISH):
                       BICYCLIC CARBOHYDRATE COMPOUNDS USEFUL IN THE TREATMENT
                        OF INFECTIONS CAUSED BY FLAVIVIRIDAE SP., SUCH AS
                        HEPATITIS C AND BOVINE VIRAL DIARRHEA
                        VIRUSES
TITLE (FRENCH):
                        COMPOSES DE CARBOHYDRATES BICYCLIQUES UTILISES DANS LE
                        TRAITEMENT D'INFECTIONS PROVOQUEES PAR LES
                        FLAVIVIRIDAE, TELS QUE LES VIRUS DE L'HEPATITE C ET LA
                        DIARRHEE VIRALE BOVINE
TITLE (GERMAN):
                        BIZYKLISCHE KOHLENHYDRATVERBINDUNGEN ZUR BEHANDLUNG VON
                        DURCH FLAVIVIRIDAE SP., WIE ETWA HEPATITIS-C- UND
                        BOVINE VIRUSDIARRHOE-VIREN AUSGELOESTEN INFEKTIONEN
INVENTOR (S):
                        SAS, Benedikt, Kwakkelstraat 71, 9190 Stekene,
                        BE; VAN HEMEL, Johan, St-Theresiastraat 44, 2600
                        Antwerpen, BE; VANDENKERCKHOVE, Jan, Vosselotstraat
                        140, 3271 Zichem, BE; PEYS, Eric, Ossenberg 81, 2490
                        Balen, BE; VAN DER EYCKEN, Johan, c/o University of
                        Ghent, Denderhoufemlaan 261, 9400 Ninove, BE;
                        RUTTENS, Bart, c/o University of Ghent, Karperstraat
                        11, 9000 Gent, BE; BALAZARINI, Jan, c/o Rega
                        Institute, Minderbroedersstraat 10, 3000 Leuven, BE;
                        DECLERCQ, Erik, c/o Rega Institute,
                        Minderbroedersstraat 10, 3000 Leuven, BE; NUYTS,
                        Johan, c/o Rega Institute, Minderbroedersstraat 10,
                        3000 Leuven, BE
PATENT APPLICANT(S):
                        Kemin Pharma Europe BVBA, 2100 Maury Street, Des
                        Moines, IA 50309, US
PATENT APPL. NUMBER:
AGENT:
                        Evans, Jacqueline Gail Victoria, et al, Greaves
                        Brewster LLP Indigo House Cheddar Business Park
                        Wedmore Road, Cheddar BS27 3EB, GB
AGENT NUMBER:
                        73005
LANGUAGE OF FILING:
                       English
LANGUAGE OF PUBL.:
                       English
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=> e SAS b/in

LANGUAGE OF PROCEDURE: English LANGUAGE OF TITLE: German; English; French DOCUMENT TYPE: Patent PATENT INFO TYPE: EPA2 Application published without search report PATENT INFORMATION: PATENT INFORMATION: KIND DATE NUMBER NUMBER KIND DATE EP 1590357 A2 20051102 WO 2004062575 20040729 AT BE BG CH CY CZ DE DK EE ES FI FR GB GR HU IE IT LI DESIGNATED STATES: LU MC NL PT RO SE SI SK TR AL LT LV MK EXTENSION STATES: EP 2003-810090 A 20031222 WO 2003-US41720 A 20031222 US 2003-438475P P 20030107 APPLICATION INFO.: PRIORITY INFO.: ANSWER 2 OF 5 PCTFULL COPYRIGHT 2006 Univentio on STN 2004062575 PCTFULL ED 20040803 EW 200431 ACCESSION NUMBER: TITLE (ENGLISH): BICYCLIC CARBOHYDRATE COMPOUNDS USEFUL IN THE TREATMENT OF INFECTIONS CAUSED BY FLAVIVIRIDAE SP., SUCH AS HEPATITIS C AND BOVINE VIRAL DIARRHEA **VIRUSES** TITLE (FRENCH): COMPOSES DE CARBOHYDRATES BICYCLIQUES UTILISES DANS LE TRAITEMENT D'INFECTIONS PROVOQUEES PAR LES FLAVIVIRIDAE, TELS QUE LES VIRUS DE L'HEPATITE C ET LA DIARRHEE VIRALE BOVINE INVENTOR(S): SAS, Benedikt, Kwakkelstraat 71, B-B-9190 Stekene, BE [BE, BE]; VAN HEMEL, Johan, St-Theresiastraat 44, B-B-2600 Antwerpen, BE [BE, BE]; VANDENKERCKHOVE, Jan, Vosselotstraat 140, B-B-3271 Zichem, BE [BE, BE]; PEYS, Eric, Ossenberg 81, B-B-2490 Balen, BE [BE, BE]; VAN DER EYCKEN, Johan, University of Ghent, Denderhoufemlaan 261, B-B-9400 Ninove, BE [BE, BE]; RUTTENS, Bart, University of Ghent, Karperstraat 11, B-B-9000 Gent, BE [BE, BE]; BALZARINI, Jan, Rega Institute, Minderbroedersstraat 10, B-B-3000 Leuven, BE [BE, BE]; DECLERCQ, Erik, Rega Institute, Minderbroedersstraat 10, B-B-3000 Leuven, BE [BE, BE]; NUYTS, Johan, Rega Institute, Minderbroedersstraat 10, B-B-3000 Leuven, BE [BE, BE] PATENT ASSIGNEE(S): KEMIN PHARMA L.C., 2100 Maury Street, Des Moines, IA 50309, US [US, US], for all designates States except US; SAS, Benedikt, Kwakkelstraat 71, B-B-9190 Stekene, BE [BE, BE], for US only; VAN HEMEL, Johan, St-Theresiastraat 44, B-B-2600 Antwerpen, BE [BE, BE], for US only; VANDENKERCKHOVE, Jan, Vosselotstraat 140, B-B-3271 Zichem, BE [BE, BE], for US only; PEYS, Eric, Ossenberg 81, B-B-2490 Balen, BE [BE, BE], for US only; VAN DER EYCKEN, Johan, University of Ghent, Denderhoufemlaan 261, B-B-9400 Ninove, BE [BE, BE], for RUTTENS, Bart, University of Ghent, Karperstraat 11, B-B-9000 Gent, BE [BE, BE], for US only; BALZARINI, Jan, Rega Institute, Minderbroedersstraat 10, B-B-3000 Leuven, BE [BE, BE], for US only; DECLERCQ, Erik, Rega Institute, Minderbroedersstraat

10, B-B-3000 Leuven, BE [BE, BE], for US only;

B-B-3000 Leuven, BE [BE, BE], for US only

NUYTS, Johan, Rega Institute, Minderbroedersstraat 10,

AGENT: HERINK, Kent\$, Davis, Brown, Koehn, Shors & Roberts, P.C., The Financial Center, 666 Walnut Street, Suite 2500, Des Moines, IA 50309\$, US LANGUAGE OF FILING: English LANGUAGE OF PUBL.: English DOCUMENT TYPE: Patent PATENT INFORMATION: NUMBER KIND DATE ------WO 2004062575 A2 20040729 DESIGNATED STATES AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR · W: CU CZ DE DK DM DZ EC EE EG ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NI NO NZ OM PG PH PL PT RO RU SC SD SE SG SK SL SY TJ TM TN TR TT TZ UA UG US UZ VC VN YU ZA ZM ZW RW (ARIPO): GH GM KE LS MW MZ SD SL SZ TZ UG ZM ZW RW (EAPO): AM AZ BY KG KZ MD RU TJ TM RW (EPO): AT BE BG CH CY CZ DE DK EE ES FI FR GB GR HU IE IT LU MC NL PT RO SE SI SK TR BF BJ CF CG CI CM GA GN GQ GW ML MR NE SN TD TG RW (OAPI): APPLICATION INFO.: WO 2003-US41720 A 20031222 PRIORITY INFO.: US 2003-60/438,475 20030107 ANSWER 3 OF 5 PCTFULL COPYRIGHT 2006 Univentio on STN ACCESSION NUMBER: 2004041176 PCTFULL ED 20040527 EW 200421 TITLE (ENGLISH): THE USE OF ENDOPEROXIDES FOR THE TREATMENT OF INFECTIONS CAUSED BY FLAVIVIRIDAE, INCLUDING HEPATITIS C, BOVINE VIRAL DIARRHEA AND CLASSICAL SWINE FEVER VIRUS TITLE (FRENCH): UTILISATION D'ENDOPEROXYDES POUR LE TRAITEMENT D'INFECTIONS CAUSEES PAR UN FLAVIVIRIDAE, TELLES OUE L'HEPATITE C, LA DIARRHEE VIRALE BOVINE ET LE VIRUS DE LA PESTE PORCINE CLASSIQUE INVENTOR(S): VANDENKERCKHOVE, Jan, Vosselotstraat 140, B-3271 Zichem, BE; SAS, Benedikt, Kwakkelstraat 71, B-9190 Stekene, BE: PEYS, Eric, Ossenberg 81, B-2490 Balen, BE; VAN HEMEL, Johan, St-Theresiastraat 44, B-2600 Antwerpen, BE PATENT ASSIGNEE(S): KEMIN PHARMA EUROPE B.V.B.A., Kristi Krafka, 2100 Maury Street, Box 70, Des Moines, IA 50301, US [US, BE] AGENT: HERINK, Kent\$, Davis, Brown, Koehn, Shors & Roberts, P.C., The Financial Center, 666 Walnut Street, Suite 2500, Des Moines, IA 50309\$, US LANGUAGE OF FILING: English LANGUAGE OF PUBL.: English DOCUMENT TYPE: Patent PATENT INFORMATION: NUMBER KIND DATE -----WO 2004041176 A2 20040521 DESIGNATED STATES W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK DM DZ EC EE EG ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NI NO NZ OM PG PH PL PT RO RU SC SD SE SG SK SL SY TJ TM TN TR TT TZ UA UG UZ VC VN YU ZA ZM ZW RW (ARIPO): GH GM KE LS MW MZ SD SL SZ TZ UG ZM ZW RW (EAPO): AM AZ BY KG KZ MD RU TJ TM AT BE BG CH CY CZ DE DK EE ES FI FR GB GR HU IE IT LU RW (EPO): MC NL PT RO SE SI SK TR

BF BJ CF CG CI CM GA GN GQ GW ML MR NE SN TD TG WO 2003-US34431 A 20031029 APPLICATION INFO.: PRIORITY INFO.: US 2002-60/422,683 20021031

RW (OAPI):

ANSWER 4 OF 5 USPATFULL on STN

ACCESSION NUMBER: 2005:69506 USPATFULL

TITLE: Use of endoperoxides for the treatment of infections

caused by flaviviridae, including hepatitis C, bovine

viral diarrhea and classical swine fever virus

INVENTOR (S): Sas, Benedikt, Stekene, BELGIUM

> hemel, Johan Van, Antwerpen, BELGIUM Vandenkerckhove, Jan, Zichem, BELGIUM

Peys, Eric, Balen, BELGIUM

NUMBER KIND DATE PATENT INFORMATION:

APPLICATION INFO.:

US 2005059647 A1 20050317

US 2003-664008 A1 20030916 (10)

DOCUMENT TYPE: FILE SEGMENT:

Utility APPLICATION

LEGAL REPRESENTATIVE:

DAVIS, BROWN, KOEHN, SHORS & ROBERTS, P.C., THE

FINANCIAL CENTER, 666 WALNUT STREET, SUITE 2500, DES

MOINES, IA, 50309-3993

NUMBER OF CLAIMS:

1

EXEMPLARY CLAIM: NUMBER OF DRAWINGS:

1 Drawing Page(s)

LINE COUNT:

355

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 5 OF 5 USPATFULL on STN

ACCESSION NUMBER:

2004:268280 USPATFULL

TITLE:

Bicyclic carbohydrate compounds useful in the treatment

of infections caused by Flaviviridae sp., such as

hepatitis C and bovine viral diarrhea

viruses

INVENTOR (S):

Sas, Benedikt, Stekene, BELGIUM

Hemel, Johan Van, Antwerpen, BELGIUM Vandenkerckhove, Jan, Zichem, BELGIUM

Peys, Eric, Balen, BELGIUM

Van der Eycken, Johan, Ninove, BELGIUM

Ruttens, Bart, Gent, BELGIUM Balzarini, Jan, Heverlee, BELGIUM Clercq, Eric De, Bierbeek, BELGIUM

Neyts, Johan, Kessel, BELGIUM

NUMBER KIND DATE -----

PATENT INFORMATION:

US 2004209823 A1 US 2003-743111 A1 20041021

APPLICATION INFO.: A1 20031222 (10)

NUMBER DATE

PRIORITY INFORMATION:

-----US 2003-438475P 20030107 (60)

DOCUMENT TYPE:

Utility

FILE SEGMENT:

APPLICATION

LEGAL REPRESENTATIVE:

DAVIS, BROWN, KOEHN, SHORS & ROBERTS, P.C., THE

FINANCIAL CENTER, 666 WALNUT STREET, SUITE 2500, DES

MOINES, IA, 50309-3993

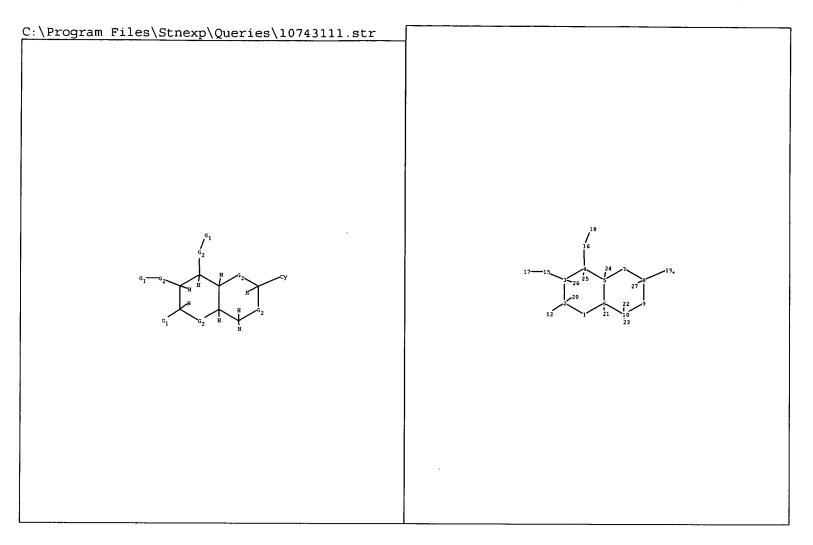
NUMBER OF CLAIMS:

EXEMPLARY CLAIM: NUMBER OF DRAWINGS:

3 Drawing Page(s)

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.



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ring nodes :

1 2 3 4 5 6 7 8 9 10

chain bonds :

2-12 2-20 3-15 3-26 4-16 4-25 5-24 6-21 8-19 8-27 10-22 10-23 15-17 16-18 ring bonds :

1-2 1-6 2-3 3-4 4-5 5-6 5-7 6-10 7-8 8-9 9-10

exact/norm bonds :

1-2 1-6 2-3 2-12 2-20 3-4 3-15 3-26 4-5 4-16 4-25 5-6 5-7 5-24 6-10 6-21 7-8 8-9 8-19 8-27 9-10 10-22 10-23 15-17 16-18

G1:Cy,Ak

G2:0,S,N

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom 12:CLASS 15:Atom 16:Atom 17:CLASS 18:CLASS 19:Atom 20:CLASS 21:CLASS 22:CLASS 23:CLASS 24:CLASS 25:CLASS 26:CLASS 27:CLASS

10/743,111

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TERMINAL (ENTER 1, 2, 3, OR ?):2

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NEWS 3 SEP 09
                 ACD predicted properties enhanced in REGISTRY/ZREGISTRY
         OCT 03
NEWS 4
                 MATHDI removed from STN
NEWS 5
         OCT 04
                 CA/CAplus-Canadian Intellectual Property Office (CIPO) added
                 to core patent offices
                 New CAS Information Use Policies Effective October 17, 2005
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         OCT 13
NEWS 7
         OCT 17
                 STN(R) AnaVist(TM), Version 1.01, allows the export/download
                 of CAplus documents for use in third-party analysis and
                 visualization tools
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NEWS 9 OCT 27 DIOGENES content streamlined
NEWS 10 OCT 27 EPFULL enhanced with additional content
NEWS 11
         NOV 14
                 CA/CAplus - Expanded coverage of German academic research
NEWS 12
         NOV 30 REGISTRY/ZREGISTRY on STN(R) enhanced with experimental
                 spectral property data
NEWS 13 DEC 05
                 CASREACT(R) - Over 10 million reactions available
NEWS 14 DEC 14 2006 MeSH terms loaded in MEDLINE/LMEDLINE
NEWS 15 DEC 14 2006 MeSH terms loaded for MEDLINE file segment of TOXCENTER
NEWS 16 DEC 14 CA/CAplus to be enhanced with updated IPC codes
        DEC 16 MARPATprev will be removed from STN on December 31, 2005
NEWS 17
NEWS 18 DEC 21 IPC search and display fields enhanced in CA/CAplus with the
                IPC reform
         DEC 23 New IPC8 SEARCH, DISPLAY, and SELECT fields in USPATFULL/USPAT2
NEWS 19
NEWS EXPRESS
            JANUARY 03 CURRENT VERSION FOR WINDOWS IS V8.01,
              CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
              AND CURRENT DISCOVER FILE IS DATED 19 DECEMBER 2005.
              V8.0 USERS CAN OBTAIN THE UPGRADE TO V8.01 AT
              http://download.cas.org/express/v8.0-Discover/
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SINCE FILE TOTAL ENTRY SESSION 0.21 0.21

FULL ESTIMATED COST

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TSCA INFORMATION NOW CURRENT THROUGH JULY 14, 2005

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REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

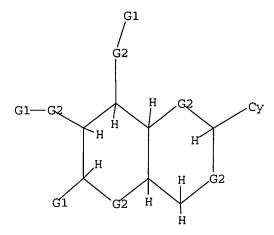
http://www.cas.org/ONLINE/UG/regprops.html

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L1 STRUCTURE UPLOADED

=> d l1 L1 HAS NO ANSWERS

L1 STR



G1 Cy,Ak G2 O,S,N

Structure attributes must be viewed using STN Express query preparation.

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1.3% PROCESSED 2000 ITERATIONS INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **INCOMPLETE**

BATCH **INCOMPLETE**

PROJECTED ITERATIONS: 3171635 TO 3218685

PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

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28.1% PROCESSED 897154 ITERATIONS

38 ANSWERS

38 ANSWERS

0 ANSWERS

31.3% PROCESSED 1000000 ITERATIONS INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)

SEARCH TIME: 00.00.21

FULL FILE PROJECTIONS: ONLINE **INCOMPLETE**

BATCH **INCOMPLETE**

PROJECTED ITERATIONS: 3196989 TO 3196989

PROJECTED ANSWERS: 3196989 TO 3196989 PROJECTED ANSWERS: 88 TO 154

L3 38 SEA SSS FUL L1

=> d 1-38 13

L3 ANSWER 1 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 867219-64-5 REGISTRY

ED Entered STN: 10 Nov 2005

CN Heptitol, 5,7-benzylidene-2,6-anhydro-1-nitro-1-desoxy-, diacetate (4CI) (CA INDEX NAME)

FS 3D CONCORD

MF C18 H21 N O9

SR CAS EARLY REGISTRATIONS

LC STN Files: CA, CAPLUS

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 2 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 863023-80-7 REGISTRY

ED Entered STN: 13 Sep 2005

CN D-glycero-D-gulo-Heptose, 2,6-anhydro-5,7-O-[(R)-phenylmethylene]-, 3,4-diacetate (9CI) (CA INDEX NAME)

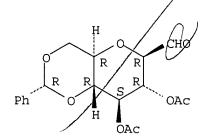
FS STEREOSEARCH

MF C18 H20 O8

SR CA

LC STN Files: CA, CAPLUS

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

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1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 3 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 863023-79-4 REGISTRY

ED Entered STN: 13 Sep 2005

CN D-Glucitol, 1,5-anhydro-1-C-(1,3-diphenyl-2-imidazolidinyl)-4,6-O-[(R)-phenylmethylene]-, 2,3-diacetate, (1S)- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C32 H34 N2 O7

SR CA

LC STN Files: CA, CAPLUS

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 4 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 863023-77-2 REGISTRY

ED Entered STN: 13 Sep 2005

CN D-glycero-D-gulo-Heptononitrile, 2,6-anhydro-5,7-O-[(R)-phenylmethylene]-, 3,4-diacetate (9CI) (CA INDEX NAME)

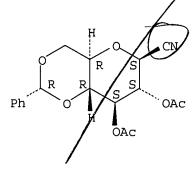
FS STEREOSEARCH

MF C18 H19 N O7

SR CA

LC STN Files: CA, CAPLUS

Absolute stereochemistry. Rotation (-).



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

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1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 5 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 863023-76-1 REGISTRY

ED Entered STN: 13 Sep 2005

CN D-glycero-D-gulo-Heptose, 2,6-anhydro-5,7-O-[(R)-phenylmethylene]-, O-acetyloxime, 3,4-diacetate (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C20 H23 N O9

SR CA

LC STN Files: CA, CAPLUS

Absolute stereochemistry.

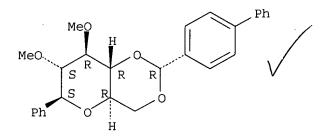
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1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

- L3 ANSWER 6 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN
- RN 850313-81-4 REGISTRY
- ED Entered STN: 12 May 2005
- CN D-Glucitol, 1,5-anhydro-4,6-O-[(R)-[1,1'-biphenyl]-4-ylmethylene]-2,3-di-O-methyl-1-C-phenyl-, (1S)- (9CI) (CA INDEX NAME)
- FS STEREOSEARCH
- MF C27 H28 O5
- SR CA
- LC STN Files: CA, CAPLUS, USPAT2, USPATFULL

Absolute stereochemistry. Rotation (-).



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 1 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- L3 ANSWER 7 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN
- RN 850313-79-0 REGISTRY
- ED Entered STN: 12 May 2005
- CN D-Glucitol, 1,5-anhydro-2,3-di-0-methyl-4,6-0-[(R)-2-

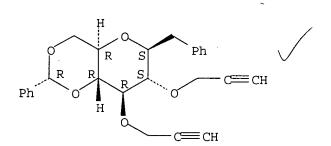
naphthalenylmethylene]-1-C-phenyl-, (1S)- (9CI) (CA INDEX NAME)

- FS STEREOSEARCH
- MF C25 H26 O5
- SR CA
- LC STN Files: CA, CAPLUS, USPAT7, USPATFULL

Absolute stereochemistry. Rotation (-).

- 1 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- L3 ANSWER 8 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN
- RN 850313-77-8 REGISTRY
- ED Entered STN: 12 May 2005
- CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-1-deoxy-1-phenyl-5,7-O-[(R)-phenylmethylene]-3,4-di-O-2-propynyl- (9CI) (CA INDEX NAME)
- FS STEREOSEARCH
- MF C26 H26 O5
- SR CA
- LC STN Files: CA, CAPLUS, USPATZ, USPATFULL

Absolute stereochemistry. Rotation (-).

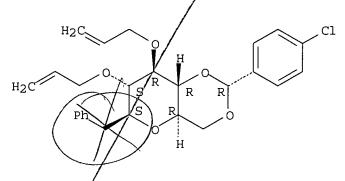


PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 1 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- L3 ANSWER 9 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN
- RN 850313-76-7 REGISTRY
- ED Entered STN: 12 May 2005
- CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-1-deoxy-1-phenyl-5,7-O-[(R)-phenylmethylene]-3,4-di-O-2-propenyl- (9CI) (CA INDEX NAME)
- FS STEREOSEARCH
- MF C26 H30 O5
- SR CA
- LC STN Files: CA, CAPLUS, USPATZ, USPATFULL

- 1 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- L3 ANSWER 10 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN
- RN 846061-34-5 REGISTRY
- ED Entered STN: 21 Mar 2005
- CN D-glycero-D-gulo-Heptitol, 2/6-anhydro-5,7-O-[(R)-(4-chlorophenyl)methylene]-1-deoxy-1-phenyl-3,4-di-O-2-propenyl- (9CI) (CA INDEX NAME)
- FS STEREOSEARCH
- MF C26 H29 Cl O5
- SR CA
- LC STN Files: CA, CAPLUS, USPATFULL

Absolute stereochemistry



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 1 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- L3 ANSWER 11 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN
- RN 846061-33-4 REGISTRY
- ED Entered STN: 21 Mar 2005
- CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-5,7-O-[(R)-(4-chlorophenyl/methylene]-1-deoxy-3,4-di-O-methyl-1-phenyl- (9CI) (CA INDEX NAME)
- FS STEREOSEARĆH
- MF C22 H25 C1 05
- SR CA
- LC STN Files: CA, CAPLUS, USPATFULL

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 12 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 846061-32-3 REGISTRY

ED Entered STN: 21 Mar 2005

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-1-deoxy-3,4-di-O-ethyl-1-phenyl-5,7-O-[(R)-phenylmethylene]- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C24 H30 O5

SR CA

LC STN Files: CA, CAPLÚS, USPAT2, USPATFULL

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1907 TO DATE)

2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 13 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 846061-31-2 REGISTRY

ED Entered STN: 21 Mar 2005

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-5,7-O-[(R)-(4-chlorophenyl)methylene]-1-deoxy-3,4-di-O-ethyl-1-phenyl- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C24 H29 Cl O5

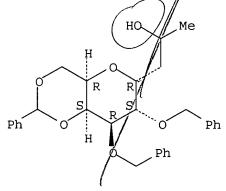
SR CA

LC STN Files: CA, CAPLUS, USPATFULL

Absolute stereochemistry. Rotation (-).

- 1 REFERENCES IN FILÉ CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- L3 ANSWER 14 OF 38 REGISTRY / COPYRIGHT 2006 ACS on STN
- RN 800394-09-6 REGISTRY
- ED Entered STN: 21 Dec 2004
- CN D-glycero-L-galacto-Nonitol, 2,6-anhydro-7,9-dideoxy-4,5-bis-O-(phenylmethyl)-1,3-O-(phenylmethylene)-, (8ξ)- (9CI) (CA INDEX NAME)
- FS STEREOSEARCH
- MF C30 H34 O6
- SR CA
- LC STN Files: CA, CAPLUS, CASREACT

Absolute stereochemistry.

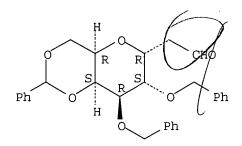


PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 1 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- L3 ANSWER 15 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN
- RN 800393-91-3 REGISTRY
- ED Entered STN: 21 Dec 2004
- CN L-glycero-L-galacto-Octitol, 2,6-anhydro-7-deoxy-4,5-bis-O-(phenylmethyl)-1,3-O-(phenylmethylene)-, acetate (9CI) (CA INDEX NAME)
- FS STEREOSEARCH
- MF C31 H34 O7
- SR CA
- LC STN Files: CA, CAPLUS, CASREACT

- 1 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- L3 ANSWER 16 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN
- RN 800393-89-9 REGISTRY
- ED Entered STN: 21 Dec 2004
- CN D-glycero-L-gluco-Octose, 3,7-anhydro-2-deoxy-4,5-bis-O-(phenylmethyl)-6,8-O-(phenylmethylene)- (9CI) (CA INDEX NAME)
- FS STEREOSEARCH
- MF C29 H30 O6
- SR CA
- LC STN Files: CA, CAPLUS, CASREACT

Absolute stereochemistry.

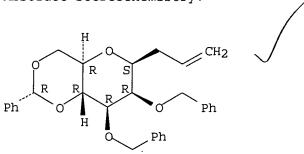


PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 1 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- L3 ANSWER 17 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN
- RN 800393-84-4 REGISTRY
- ED Entered STN: 21 Dec 2004
- CN D-glycero-L-galacto-Non-8-enitol, 2,6-anhydro-7,8,9-trideoxy-4,5-bis-0-(phenylmethyl)-1,3-O-(phenylmethylene)- (9CI) (CA INDEX NAME)
- FS STEREOSEARCH
- MF C30 H32 O5
- SR CA
- LC STN Files: CA, CAPLUS, CASREACT

- 1 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- L3 ANSWER 18 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN
- RN 795306-04-6 REGISTRY
- ED Entered STN: 09 Dec 2004
- CN D-glycero-D-galacto-Non-1-enitol, 4,8-anhydro-1,2,3-trideoxy-5,6-bis-0-(phenylmethyl)-7,9-O-[(R)-phenylmethylene]- (9CI) (CA INDEX NAME)
- FS STEREOSEARCH
- MF C30 H32 O5
- SR CA
- LC STN Files: CA, CAPLUS, CASREACT

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 1 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- L3 ANSWER 19 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN
- RN 770727-53-2 REGISTRY
- ED Entered STN: 28 Oct 2004
- CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-5-O-[2,3-bis-O-(phenylmethyl)-4,6-O-[(R)-phenylmethylene]- β -D-glucopyranosyl]-4,7-bis-O-(phenylmethyl)-1,3-O-[(R)-phenylmethylene]- (9CI) (CA INDEX NAME)
- FS STEREOSEARCH
- MF C55 H56 O11
- SR CA
- LC STN Files: CA, CAPLUS

Absolute stereochemistry. Rotation (+).

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 20 OF 38 REGISŢŔY COPYRIGHT 2006 ACS on STN

RN 770727-51-0 REGISTRY

ED Entered STN: 28 Oct 2004

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-5-0-[2,3-di-O-acetyl-4,6-0-[(R)-phenylmethylene]-β-D-glucopyranosyl]-1,3-0-[(R)-phenylmethylene]-, diacetate (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C35 H40 O15

SR CA

LC STN Files: CA, CAPLUS

Absolute stereochemistry. Rotation (+).

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 1 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 21 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN RN 727416-80-0 REGISTRY

ED Entered STN: 16 Aug 2004

CN D-Glucitol, 1,5-anhydro-2,3-di-0-methyl-1-C-phenyl-4,6-0-[(R)-[4-(trifluoromethyl)phenyl]methylene]-, (1S)- (9CI) (CA INDEX NAME)

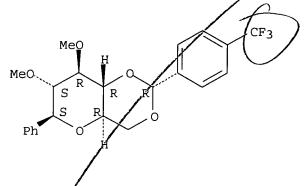
FS STEREOSEARCH

MF C22 H23 F3 O5

SR CA

LC STN Files: CA, CAPLUS, TOXQENTER, USPATFULL

Absolute stereochemistry. Rotation (-).

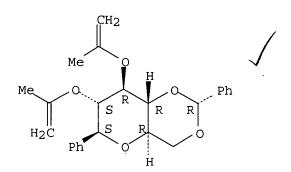


PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

- L3 ANSWER 22 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN
- RN 727415-96-5 REGISTRY
- ED Entered STN: 16 Aug 2004
- CN D-Glucitol, 1,5-anhydro-2,3-bis-O-(1-methylethenyl)-1-C-phenyl-4,6-O-[(R)-phenylmethylene]-, (1S)- (9CI) (CA INDEX NAME)
- FS STEREOSEARCH
- MF C25 H28 O5
- SR CA
- LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

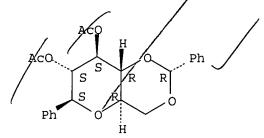
- 1 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- L3 ANSWER 23 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN
- RN 727415-94-3 REGISTRY
- ED Entered STN: 16 Aug 2004
- CN D-Glucitol, 1,5-anhydro-1-C-phenyl-4,6-O-[(R)-phenylmethylene]-, diacetate, (1S)- (9CI) (CA INDEX NAME)

FS STEREOSEARCH MF C23 H24 O7

SR CA

LC STN Files: CA, ÇAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry. Rotation (-).



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 24 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 727415-92-1 REGISTRY

ED Entered STN: 16 Aug 2004

CN D-Glucitol, 1,5-anhydro-2,3-bis-O-(1-methylethyl)-1-C-phenyl-4,6-O-[(R)-phenylmethylene]-, (1S)- (9CI) (CA INDEX NAME)

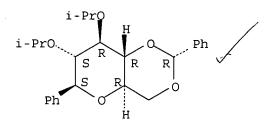
FS STEREOSEARCH

MF C25 H32 O5

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry. Rotation (-).



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 25 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 727415-91-0 REGISTRY

ED Entered STN: 16 Aug 2004

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-3,4-di-O-butyl-1-deoxy-1-phenyl-5,7-O-[(R)-phenylmethylene]- (9CI) (CA INDEX NAME)

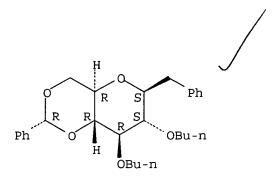
FS STEREOSEARCH

MF C28 H38 O5

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry. Rotation (-).



1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 26 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 727415-90-9 REGISTRY

ED Entered STN: 16 Aug 2004

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-1-deoxy-1-phenyl-5,7-O-[(R)-phenylmethylene]-3,4-di-O-propyl- (9CI) (CA INDEX NAME)

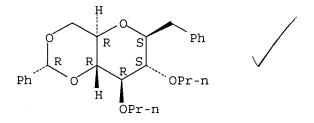
FS STEREOSEARCH

MF C26 H34 O5

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry. Rotation (-).



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 27 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 648431-84-9 REGISTRY

ED Entered STN: 10 Feb 2004

CN D-glycero-L-gulo-Heptitol, 5-(acetylamino)-2,6-anhydro-5,7-dideoxy-4-0-(2,3-di-0-benzoyl-6-deoxy-4-0-methyl- α -L-mannopyranosyl)-7-

(diethoxyphosphinyl)-1,3-0-[(R)-phenylmethylene]- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C41 H50 N O14 P

SR CA

LC STN Files: CA, CAPLUS

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 28 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 648431-79-2 REGISTRY

ED Entered STN: 10 Feb 2004

CN D-glycero-D-ido-Heptonic acid, 3-(acetylamino)-2,6-anhydro-3-deoxy-4-0-(2,3-di-0-benzoyl-6-deoxy-4-0-methyl-α-L-mannopyranosyl)-5,7-0-[(R)-phenylmethylene]-, methyl ester (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C38 H41 N O13

SR CA

LC STN Files: CA, CAPLUS

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 1 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 29 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 648431-78-1 REGISTRY

ED Entered STN: 10 Feb 2004

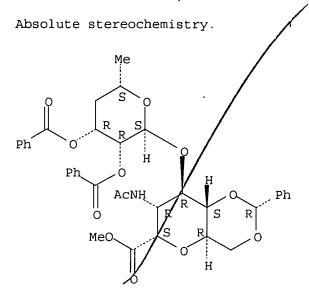
CN D-glycero-D-ido-Heptonic acid, 3-(acetylamino)-2,6-anhydro-3-deoxy-4-0- $(2,3-di-0-benzoyl-4,6-dideoxy-\alpha-L-lyxo-hexopyranosyl)-5,7-O-[(R)-phenylmethylene]-, methyl ester (9CI) (CA INDEX NAME)$

FS STEREOSEARCH

MF C37 H39 N O12

SR CA

LC STN Files: CA, CAPLUS



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 30 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 623148-59-4 REGISTRY

ED Entered STN: 03 Dec 2003

CN D-glycero-L-gulo-Heptitol, 2,6-anhydro-4,5-bis-O-[(4-methoxyphenyl)methyl]-7-C-phenyl-1,3-O-[(R)-phenylmethylene]-7-O-[[2-(trimethylsilyl)ethoxy]methyl]-, (7R)- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C42 H52 O9 Si

SR CA

LC STN Files: CA, CAPLUS, CASREACT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 31 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 623148-58-3 REGISTRY

ED Entered STN: 03 Dec 2003

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-3,4-bis-O-[(4-methoxyphenyl)methyl]-1-C-phenyl-5,7-O-[(R)-phenylmethylene]-, (1S)- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C36 H38 O8

SR CA

LC STN Files: CA, CAPLUS, CASREACT

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 1 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 32 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN RN 623148-57-2 REGISTRY ED Entered STN: 03 Dec 2003 D-glycero-L-gulo-Heptitol, 2,6-anhydro-4,5-bis-O-[(4-methoxyphenyl)methyl]-7-C-phenyl-1,3-O-[(R)-phenylmethylene]-, (7R)- (9CI) (CA INDEX NAME) CN FS STEREOSEARCH C36 H38 O8 MF SR CA CA, CAPLUS, CASREACT LC STN Files: Absolute stereochemistry. ØМе R R R Η Ph

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 33 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 592532-99-5 REGISTRY

ED Entered STN: 25 Sep 2003

CN D-glycero-L-galacto-Non-8-enitol, 2,6-anhydro-7,8,9-trideoxy-4,5-bis-0-(phenylmethyl)-1,3-O-[(S)-phenylmethylene]- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C30 H32 O5

SR CA

LC STN Files: CA, CAPLUS, CASREACT

1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 34 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 511274-46-7 REGISTRY

ED Entered STN: 06 May 2003

CN D-glycero-D-gulo-Non-1-enitol, 4,8-anhydro-1,2,3-trideoxy-5,6-di-O-methyl-7,9-O-[(R)-phenylmethylene]- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H24 O5

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 35 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN

RN 511274-42-3 REGISTRY

ED Entered STN: 06 May 2003

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-1-deoxy-3,4-di-O-methyl-1-phenyl-5,7-O-[(R)-phenylmethylene]- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C22 H26 O5

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATZ, USPATFULL

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3 REFERENCES IN FILE CA (1907 TO DATE)

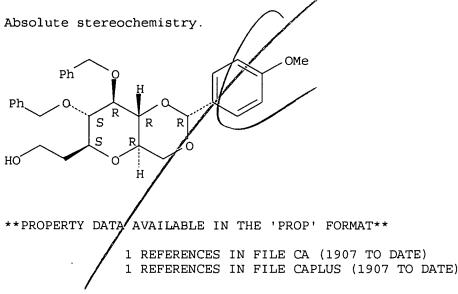
3 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L3 ANSWER 36 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN RN 511274-41-2 REGISTRY

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Entered STN: 06 May 2003
ED
     D-Glucitol, 1,5-anhydro-2,3-di-O-methyl-1-C-phenyl-4,6-O-[(R)-
CN
     phenylmethylene]-, (1S)- (9CI) (CA INDEX NAME)
FS
     STEREOSEARCH
     C21 H24 O5
MF
SR
     CA
     STN Files:
                  CA, CAPLUS, TOXCENTER, USPATZ, USPATFULL
LC
Absolute stereochemistry. Rotation (-).
      MeQ
                    Ph
MeQ.
      S
            R
 Ph
**PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**
               3 REFERENCES IN FILE CA (1907 TÓ DATE)
               3 REFERENCES IN FILE CAPLUS (1907 TO DATE)
L3
     ANSWER 37 OF 38 REGISTRY
                                COPYRIGHT 2006 ACS on STN
     482646-98-0 REGISTRY
RN
ED
     Entered STN: 29 Jan 2003
CN
     D-glycero-D-gulo-Octitol, 3,7-anhydro-2-deoxy-6,8-0-[(R)-(4-
     methoxyphenyl)methylene]-1,4,5/tris-O-(phenylmethyl)- (9CI) (CA INDEX
     STEREOSEARCH
FS
MF
     C37 H40 O7
SR
                  CA, CAPLUS
                               CASREACT, TOXCENTER
LC
     STN Files:
Absolute stereochemistry.
                                         OMe
                 S
                       R
Ph
                      H
**PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**
               1 REFERENCES IN FILE CA (1907 TO DATE)
               1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
L3
     ANSWER 38 OF 38 REGISTRY COPYRIGHT 2006 ACS on STN
     482646-97-9 REGISTRY
RN
ED
     Entered STN: 29 Jan 2003
     D-glycero-D-gulo-Octitol, 3,7-anhydro-2-deoxy-6,8-O-[(R)-(4-
CN
     methoxyphenyl) methylene] -4,5-bis-O-(phenylmethyl) - (9CI) (CA INDEX NAME)
FS
     STEREOSEARCH
MF
     C30 H34 O7
```

SR CA

LC STN Files: CA, CAPLUS, CASREACT, TOXCENTER



=> file caplus
COST IN U.S. DOLLARS

FULL ESTIMATED COST

SINCE FILE TOTAL ENTRY SESSION 239.58 239.79

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=> d his

(FILE 'HOME' ENTERED AT 20:01:38 ON 06 JAN 2006)

FILE 'REGISTRY' ENTERED AT 20:01:48 ON 06 JAN 2006

L1 STRUCTURE UPLOADED

L2 0 S L1 SAM L3 38 S L1 FULL

FILE 'CAPLUS' ENTERED AT 20:03:10 ON 06 JAN 2006

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=> d 13
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Absolute stereochemistry.

YOU HAVE REQUESTED DATA FROM FILE 'REGISTRY' - CONTINUE? (Y) /N:n

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=> s 13
            15 L3
L4
=> d l4 bib abs hitstr 1-15
     ANSWER 1 OF 15 CAPLUS
                             COPYRIGHT 2006 ACS on STN
ΑN
     2005:1146690 CAPLUS
DN
     144:23075
TΙ
     A Convergent Ring-Closing Met/athesis Approach to Carbohydrate-Based
     Macrolides with Potential Antibiotic Activity
AU
     Blom, Petra; Ruttens, Bart; Van Hoof, Steven; Hubrecht, Idzi; Van der
     Eycken, Johan; Sas, Benedikt; Van hemel, Johan; Vandenkerckhove, Jan
     Laboratory for Organic and/Bioorganic Synthesis, Department of Organic
CS
     Chemistry, Ghent University, Ghent, B-9000, Belg.
     Journal of Organic Chemistry (2005), 70(24), 10109-10112
SO
     CODEN: JOCEAH; ISSN: 0022-3263
PB
     American Chemical Societ
DT
     Journal
LΑ
     English
AB
     An efficient convergent approach has been developed for the construction
     of novel, non-natural,/carbohydrate-based macrolides. The key step in the
     synthesis is the formation of the macrocyclic ring via a ring-closing
     metathesis reaction. / The obtained macrolide analogs have been screened
     for biol. activity against Gram-pos. and Gram-neg. bacteria, including
     resistant strains, yeasts, and molds.
IT
     511274-41-2P 511274/42-3P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (convergent ring-closing metathesis approach to carbohydrate-based
        macrolides with potential antibiotic activity)
RN
     511274-41-2 CAPLUS
CN
     D-Glucitol, 1,5-4nhydro-2,3-di-O-methyl-1-C-phenyl-4,6-O-[(R)-
     phenylmethylene]/-, (1S) - (9CI) (CA INDEX NAME)
Absolute stereochemistry. Rotation (-).
      MeQ
MeQ.
                    Ph
      S
            R
           H
RN
     511274-42/3 CAPLUS
CN
     D-glycerd-D-gulo-Heptitol, 2,6-anhydro-1-deoxy-3,4-di-O-methyl-1-phenyl-
     5,7-0-[(元)-phenylmethylene]- (9CI) (CA INDEX NAME)
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OMe
             ON
RE.CNT
        36
              THERE ARE 36 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
L4
     ANSWER 2 OF 15 CAPLUS COPYRIGHT 2006 ACS on STN
ΑN
     2005:560681 CAPLUS
DN
     143:230118
TI
     Selective Reductions of C-(4/6-O-Benzylidene-β-D-qlucopyranosyl)
     Simo, Ondrej; Michael, Katja; Yoshida, Wesley; Chertkov, Vyacheslav;
AU
     Gross, Paul
     Department of Chemistry, University of the Pacific, Stockton, CA, USA
CS
     Synthetic Communications (2005) 35(12), 1589-1599
SO
     CODEN: SYNCAV; ISSN: 003 7-7911
PB
     Taylor & Francis, Inc.
DT
     Journal
LΑ
     English
     The nitromethyl group of C-(4,6-0-benzylidene-ss-D-glucopyranosyl)
AΒ
     nitromethane was manifulated by various reduction and oxidation methods and
     further functionalizations into -CH2NHOH, -CH=NOH, -CN, -CH=O, and
     -CH2NHCONH2, all with retention of the 4,6-0-benzylidene group. Certain
     reduction methods gave rise to a novel secondary amine via an unusual dimeric
     aminal.
IT
     863023-76-1P 86302/3-79-4P
     RL: RCT (Reactant /; SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (chemoselecti fe redns. of the nitromethyl group of C-(4,6-0-benzylidene-
        ss-D-glucopymanosyl) nitromethane containing glycosides and
        oligosaccharides)
     863023-76-1 CAPLUS
RN
     D-glycero-D-gulo-Heptose, 2,6-anhydro-5,7-O-[(R)-phenylmethylene]-,
CN
     O-acetyloxime 3,4-diacetate (9CI) (CA INDEX NAME)
Absolute stereochemistry.
Double bond geometry unknown.
          R
                   OAc
Ph
         Η
             OAc
RN
     863023/79-4 CAPLUS
```

CN D-Gluditol, 1,5-anhydro-1-C-(1,3-diphenyl-2-imidazolidinyl)-4,6-O-[(R)-phenylmethylene]-, 2,3-diacetate, (1S)- (9CI) (CA INDEX NAME)

IT 863023-77-2P 863023-80-7P

RL: SPN (Synthetic preparation); PREP (Preparation) (chemoselective redns. of the nitromethyl group of C-(4,6-0-benzylideness-D-glucopyranosyl) nitromethane containing glycosides and oligosaccharides)

RN 863023-77-2 CAPLUS

CN D-glycero-D-gulo-Heptononitrile, 2,6-anhydro-5,7-O-[(R)-phenylmethylene]-, 3,4-diacetate (9¢I) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

RN 863023-80 7 CAPLUS

CN D-glycero-D-gulo-Heptose, 2,6-anhydro-5,7-O-[(R)-phenylmethylene]-, 3,4-diacetate (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RE.CNT 20

THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L4 ANSWER 3 OF 15 CAPLUS COPYRIGHT 2006 ACS on STN
- AN 2005:371015 CAPLUS
- DN 142:411583
- TI Bicyclic carbohydrate compounds useful in the treatment of infections caused by Herpesviridae
- IN Sas, Benedikt; Van Hemel, Johan; Vandenkerckhove, Jan; Peys, Eric; Van Der Eycken, Johan; Ruttens, Bart; Blom, Petra
- PA Belg.
- SO U.S. Pat. Appl. Publ., 18 pp. CODEN: USXXCO

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DT
     Patent
LΑ
     English
FAN.CNT 2
```

| • | 2 114 | | | | | | | | | | | | | | | | | | |
|---|-----------------|-------------------|------|-----|-----|-------------|------|----------|---------------------|-----------------|-----------------|-----|-----|--------|-----|----------|-----|-----|-----|
| | | PATENT NO. | | | | KIND | | DATE | | i | APPLICATION NO. | | | | | DATE | | | |
| Р | ·Ι | US 2005090452 | | | | A1 | | 20050428 | | 1 | US 2003-690914 | | | 14 | | 20031022 | | | |
| | | US 2005267048 | | | | A9 | | 20051201 | | | | | | | | | | | |
| | | US 2003158243 | | | | A1 | | 20030821 | | US 2001-977478 | | | | | | 20011015 | | | |
| | | US 2004224904 - | | | | A1 2004 | | | 1111 US 2004-861768 | | | | | | | 20040604 | | | |
| | | WO 2005062748 | | | | A2 20050714 | | | | WO 2004-US34886 | | | | | | 20041022 | | | |
| | | | W: | ΑE, | AG, | АL, | AM, | AT, | AU, | ΑZ, | BA, | BB, | ВG, | BR, | BW, | BY, | ΒZ, | CA, | CH, |
| | | | | CN, | CO, | CR, | CU, | CZ, | DE, | DK, | DM, | DZ, | EC, | EE, | EG, | ES, | FΙ, | GB, | GD, |
| | | | | | | | | | ID, | | | | | | - | • | | | |
| | | | | | | | | | LV, | | | | | | | | | | |
| | | | | NO, | ΝŻ, | OM, | PG, | PH, | PL, | PT, | RO, | RU, | SC, | SD, | SE, | SG, | SK, | SL, | SY, |
| | | | | | - | | • | | TZ, | | | - | | | • | • | | • | |
| | | | RW: | | | | | | MW, | | | | | | | | | | |
| | | | | | | | | | RU, | | | | | | | | | | |
| | | | | | | | | | GR, | | | | | | - | - | - | - | |
| | | | | | | | BF, | BJ, | CF, | CG, | CI, | CM, | GA, | GN, | GQ, | GW, | ML, | MR, | NE, |
| ъ | דעם | пс | 2001 | • | TD, | | 7. 0 | | 2001 | 1015 | | | | | | | | | |
| P | KAI | AI US 2001-977478 | | | | | | | | | | | | | | | | | |
| | IIS 2003-690914 | | | | | Δ | | 2003 | しいノノ | | | | | | | ١. | | | |

US 2003-690914 20031022

OS MARPAT 142:411583

GI

$$R^2X$$
 XR^3
 XR^4
 XR^5
 XR^5

ΙI

AB Bicyclic glycoside I, wherein wherein R1 is either Bn, Ph; R2 and R3 are either alkyl, aryl, allyl, H; R4 and R5 form a ring and are either CH(Ph) or CH(aryl); X is either O, N or S, were prepared and tested for the treatment of infections caused by Herpseviridae, and in particular cytomegalovirus. The compds. were screened against various pathogenic viruses and more specific the human Cytomegalovirus (CMV). Antiviral activity of these new mols. are in the same range as the activity of ganciclovir for CMV screenings. Thus, bicyclic glycoside II was prepared and tested in vitro for the treatment of infections caused by Cytomegalovirus (EC50 = 0.47 to 3.2 μ g/mL).

ΙT 511274-41-2P 511274-42-3P 846061-32-3P 850313-76-7P 850313-77-8P 850313-79-0P 850313-81-4P

> RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of bicyclic glycosides as antiviral agents useful in treatment of infections caused by herpesviridae)

RN 511274-41-2 CAPLUS

CN D-Glucitol, 1,5-anhydro-2,3-di-O-methyl-1-C-phenyl-4,6-O-[(R)phenylmethylene] -, (1S) - (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

RN 511274-42-3 CAPLUS

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-1-deoxy-3,4-di-O-methyl-1-phenyl-5,7-O-[(R)-phenylmethylene]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 846061-32-3 CAPLUS

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-1-deoxy-3,4-di-O-ethyl-1-phenyl-5,7-O-[(R)-phenylmethylene]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 850313-76-7 CAPLUS

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-1-deoxy-1-phenyl-5,7-0-[(R)-phenylmethylene]-3,4-di-O-2-propenyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 850313-77-8 CAPLUS

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-1-deoxy-1-phenyl-5,7-O-[(R)-phenylmethylene]-3,4-di-O-2-propynyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

RN 850313-79-0 CAPLUS

CN D-Glucitol, 1,5-anhydro-2,3-di-O-methyl-4,6-O-[(R)-2-naphthalenylmethylene]-1-C-phenyl-, (1S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

RN 850313-81-4 CAPLUS

CN D-Glucitol, 1,5-anhydro-4,6-O-[(R)-[1,1'-biphenyl]-4-ylmethylene]-2,3-di-O-methyl-1-C-phenyl-, (1S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

- L4 ANSWER 4 OF 15 CAPLUS COPYRIGHT 2006 ACS on STN
- AN 2005:238681 CAPLUS
- DN 142:261733
- TI Preparation of bicyclic glycosides as antiviral agents for the treatment of infections caused by the Alphaherpesvirinae HSV-1 and HSV-2
- IN Sas, Benedikt; Van Hemel, Johan; Vandenkerckhove, Jan; Peys, Eric; Van Der Eycken, Johan; Van Hoof, Steven
- PA Belq.
- SO U.S. Pat. Appl. Publ., 15 pp. CODEN: USXXCO
- DT Patent
- LA English
- FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

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PΙ
      US 2005059612
                                      20050317
                                                     US 2003-663962
                                                                                 20030916
                               Α1
      WO 2005048921
                                                     WO 2004-US30205
                                                                                 20040916
                               A2
                                      20050602
      WO 2005048921
                               А3
                                      20051208
              AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
               CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,
               NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY,
               TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
          RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM,
               AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK,
               EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE,
               SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,
               SN, TD, TG
PRAI US 2003-663962
                                      20030916
                               Α
```

OS MARPAT 142:261733

GI

$$\begin{array}{c} X^2Et \\ EtX^2 \\ Y^1 \\ Y^2 \\ \end{array}$$

relatives /

Novel bicyclic glycosides I, wherein X1-X3 and Y1 and Y2 are independently AΒ selected from the group consisting of O, N, and S; Z is selected from the group consisting of F, Cl, and Br, as well as analogs, prodrugs and pharmaceutically acceptable salts thereof, together with pharmaceutical compns. for the prophylaxis and treatment of diseases caused by infections of Alphaherpesvirinae and are effective for the prophylaxis and treatment of diseases caused by infections of the Alphaherpesvirinae HSV-1 and HSV-2. Thus, glycoside II was prepared and tested as antiviral agent against HSV-1 (EC50 = 3 μ g/mL) and HSV-2 (EC50 < 0.03 μ g/mL) viruses. The mols. were screened in vitro against a series of viruses such as West Nile virus, human cytomegalo virus (HCMV), herpes simplex virus type 1 (HSV-1), herpes simplex virus type 2 (HSV-2) and varicella zoster virus (VZV).

IT 846061-31-2P

> RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES

(preparation of bicyclic glycosides as antiviral agents for the treatment of infections caused by the Alphaherpesvirinae HSV-1 and HSV-2)

RN 846061-31-2 CAPLUS

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-5,7-0-[(R)-(4chlorophenyl)methylene]-1-deoxy-3,4-di-O-ethyl-1-phenyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

IT 846061-32-3 846061-33-4 846061-34-5

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(preparation of bicyclic glycosides as antiviral agents for the treatment of infections caused by the Alphaherpesvirinae HSV-1 and HSV-2)

RN 846061-32-3 CAPLUS

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-1-deoxy-3,4-di-O-ethyl-1-phenyl-5,7-O-[(R)-phenylmethylene]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 846061-33-4 CAPLUS

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-5,7-0-[(R)-(4-chlorophenyl)methylene]-1-deoxy-3,4-di-O-methyl-1-phenyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 846061-34-5 CAPLUS

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-5,7-0-[(R)-(4chlorophenyl)methylene]-1-deoxy-1-phenyl-3,4-di-0-2-propenyl- (9CI) (CA INDEX NAME)

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S
                S
                     Η
L4
     ANSWER 5 OF 15 CAPLUS COPYRIGHT 2006 ACS on STN
     2004:830839 CAPLUS
ΑN
DN
     142:23422
TI
     Intramolecular aldol cyclization of C-4-ulopyranosyl-2'-oxoalkanes
     controlled by steric effects. Asymmetric synthesis of substituted
     8-oxabicyclo[3.2.1] octanones and -octenones and cyclopentenones
ΑU
     Zou, Wei; Shao, Huawu; Wu, Shith-Hsiung
     Institute for Biological Sciences, National Research Council of Canada,
CS
     Ottawa, ON, K1A OR6, Can.
SO
     Carbohydrate Research (2004), 339(15), 2475-2485
     CODEN: CRBRAT; ISSN: 0008-$215
PB
     Elsevier B.V.
DT
     Journal
LΑ
     English
OS
     CASREACT 142:23422
     Whereas C-2- and 4-ulopyranosyl compds. (C-2- and C-4-ulosides) can be
AΒ
     converted to cyclopentenones under base conditions through
     \beta-elimination and rin\phi contraction, base-initiated \beta-elimination
     of C-glycosyl 2'-ald#hydes and 2'-ketones results in the formation of
     acyclic \alpha, \beta-unsatd. Aldehydes or ketones. By combining both
     mol. features, 1-C-/(4-ulopyranosyl)-2-oxoalkanes were synthesized and
     their reactions when treated with base were investigated. Both \alpha-
     and \beta-anomers of q-(4-ulopyranosyl)acetaldehydes underwent a fast
     intramol. aldol reaction between the C-5 enolate and 2'-aldehyde to form
     optically pure 8 foxabicyclo[3.2.1] octanones, which further transformed to
     8-oxabicyclo[3.2/.1]octenones by eta-elimination. However, this aldol
     reaction did not occur when 1-C-(4-ulopyranosyl)propan-2-one was treated
     with base because of steric hindrance exerted by the addnl. Me group.
     Instead, an alternate C-3 enolization led to \beta-elimination and
     further elect#o-ring opening to form an acyclic enol, which was then
     converted through a disrotatory intramol. aldol cyclization to a
     cis-substituted cyclopentenone.
IT
     800393-84-4P 800393-89-9P 800393-91-3P
     800394-09-61
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (asym. synthesis of substituted 8-oxabicyclo[3.2.1]octanones and
        -octen\phines and cyclopentenones via intramol. aldol cyclization,
        \beta-elimination, and electro-ring opening)
RN
     800393-84-4 CAPLUS
```

D-glyceró-L-galacto-Non-8-enitol, 2,6-anhydro-7,8,9-trideoxy-4,5-bis-0-

(phenylmethyl) -1,3-O-(phenylmethylene) - (9CI) (CA INDEX NAME)

Absolute stereochemistry.

CN

RN 800393-89-9 CAPLUS

CN D-glycero-L-gluco-Octose, 3,7-anhydro-2-deoxy-4,5-bis-O-(phenylmethyl)-6,8-O-(phenylmethylene)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 800393-91-3 CAPLUS

CN L-glycero-L-ga/acto-Octitol, 2,6-anhydro-7-deoxy-4,5-bis-O-(phenylmethyl)-1,3-O-(phenylmethylene)-, acetate (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 800394-09-6 CAPLUS

CN D-glycer -L-galacto-Nonitol, 2,6-anhydro-7,9-dideoxy-4,5-bis-0-(phenylmethyl)-1,3-0-(phenylmethylene)-, (8ξ)- (9CI) (CA INDEX NAME)

RE.CNT 50 THERE ARE 50 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 6 OF 15 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2004:773147 CAPLUS

DN 141:424352

TI An Efficient Synthesis of β -C-Glycosides Based on the Conformational Restriction Strategy: Lewis Acid Promoted Silane Reduction of the Anomeric Position with Complete Stereoselectivity

AU Terauchi, Masaru; Abe, Hiroshi; Matguda, Akira; Shuto, Satoshi

CS Graduate School of Pharmaceutical Sciences, Hokkaido University, Sapporo, 060-0812, Japan

SO Organic Letters (2004), 6(21), 3/51-3754 CODEN: ORLEF7; ISSN: 1523-7060

Ι

PB American Chemical Society

DT Journal

LA English

OS CASREACT 141:424352

GΙ

BnO O Me

The reduction of glycono-lactols having an anomeric carbon substituent by Et3SiH/TMSOTf proceeded with complete stereoselectivity to produce the corresponding $\beta\text{-C-glycosides}$, e.g. I, when the substrates were conformationally restricted in the 4Cl-chair form by a 3,4-0-cyclic di-ketal or a 4,6-0-benzylidene protecting group. Thus, the efficient construction of $\beta\text{-C-glycosides}$ was achieved on the basis of the conformation restriction strategy.

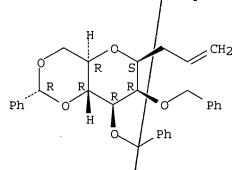
IT 795306-04-6P

RL: SPN (Synthetic preparation); PREP (Preparation) (synthesis of β-C-glycosides based on conformational restriction strategy and stereoselective Lewis acid promoted silane reduction of anomeric position)

RN 795306-04-6 CAPLUS

CN D-glycero-D-galacto-Non-1-enitol, 4,8-anhydro-1,2,3-trideoxy-5,6-bis-0-(phenylmethyl) 7,9-0-[(R)-phenylmethylene] - (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RE.CNT 37 THERE ARE 37 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 7 OF 15 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2004:610047 CAPLUS

- DN 141:134051
- TI Bicyclic carbohydrates as antiprotozoal bioactive for the treatment of infections caused by parasites
- IN Sas, Benedikt; Van Hemel, Johan; Vandenkerckhove, Jan; Van Hemel, Johan; Peys, Eric; Van Der Eycken, Johan; Ruttens, Bart; Van Hoof, Steven
- PA Kemin Pharma Europe B.V.B.A., USA
- SO PCT Int. Appl., 26 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN CNT 1

| FAN. | CNT. | T | | | | | | | | | | | | | | | | |
|------|----------------------|------------|-------|------|-----|----------|-----|------|------|-----|------|------|-------|--------|-----|-----|------|-----|
| | PAT | ENT 1 | NO. | | | KIN | D | DATE | | i | APPL | ICAT | ION 1 | NO. | | D | ATE | |
| | | | | | | | - | | | | | | | | | | | |
| ΡI | WO : | 2004 | 0625 | 90 | | A2 | | 2004 | 0729 | Ţ | WO 2 | 004- | US31 | 1 | | 2 | 0040 | 107 |
| | WO 2004062590 | | | A3 | | 20050407 | | | | | | | | | | | | |
| | | W : | ΑE, | ΑE, | AG, | AL, | AL, | AM, | AM, | AM, | ΑT, | ΑT, | AU, | AU, | AZ, | ΑZ, | BA, | BB, |
| | | | BG, | BG, | BR, | BR, | BW, | BY, | ΒY, | ΒZ, | ΒZ, | CA, | CH, | CN, | CN, | CO, | CO, | CR, |
| | | | CR, | CU, | CU, | CZ, | CZ, | DE, | DE, | DK, | DK, | DM, | DZ, | EC, | EC, | EE, | EE, | EG, |
| | | | ES, | ES, | FI, | FI, | GB, | GD, | GE, | GE, | GH, | GH, | GH, | GM, | HR, | HR, | HU, | HU, |
| | | | ID, | IL, | IN, | IS, | JP, | JΡ, | ΚE, | ΚE, | KG, | KG, | ΚP, | ΚP, | ΚP, | KR, | KR, | ΚZ, |
| | | | ΚZ, | ΚZ, | LC, | LK, | LR, | LS, | LS, | LT, | LU, | LV, | MA, | MD, | MD, | MG, | MK, | MN, |
| | | | MW, | MX, | MX, | MZ | | | | | | | | | | | | |
| (| | | 1808. | | | | | 2004 | 0916 | Ī | JS 2 | 004- | 7527 | 92 | 1 | 20 | 0040 | 107 |
| PRAI | <u>US</u> . | 2003 | -438 | 474P | | P | | 2003 | 0107 | | | | 0 | ما ـ ١ | کی | | | |
| OS | OS MARPAT 141:134051 | | | | | | | | | | | | A1 | MY & | / — | | | |

AB The use of bicyclic carbohydrates for the treatment of parasite infections is described. Different bicyclic carbohydrates have been tested in vitro against a number of protozoa. These compds. also have been screened against viruses, tumors, bacteria and fungi. Compound Al, a thiophenyl-containing bicyclic carbohydrate possessed significant activity against Trypanosoma

brucei rhodesiense, a parasite that causes the lethal sleeping sickness.
Compound A2 and Compound A3, bicyclic carbohydrates with halogen containing

groups, possessed significant activity against Leishmania donovani, a parasite that causes leishmaniasis. Bicyclic carbohydrates in general, and Compound A1, Compound A2 and Compound A3 more specifically, could be possible treatments for the sleeping sickness and leishmaniasis in the future.

IT 727416-80-0P

L4

RL: BSU (Biological study, unclassified); PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(bicyclic carbohydrates as antiprotozoal agent for treatment of parasite infections)

RN 727416-80-0 CAPLUS

CN D-Glucitol, 1,5-anhydro-2,3-di-0-methyl-1-C-phenyl-4,6-0-[(R)-[4-(trifluoromethyl)phenyl]methylene]-, (1S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

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AN
     2004:610040 CAPLUS
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DN 141:134050

ΤI Bicyclic carbohydrate compounds useful in the treatment of infections caused by Flaviviridae sp., such as hepatitis C and bovine viral diarrhea

Sas, Benedikt; Van Hemel, Johan; Vandenkerckhove, Jan; Peys, Eric; Van Der IN Eycken, Johan; Ruttens, Bart; Balzarini, Jan; Declercq, Erik; Nuyts, Johan

PA Kemin Pharma L.C., USA

SO PCT Int. Appl., 23 pp.

CODEN: PIXXD2

DTPatent

LΑ English

| FAN. | CNT | 1 | | | | | | | | | | | | | | | | |
|------|-----|--------|-------|------|-----|-----|-----|------|------|-----|------|-------|-------|-----|-----|-----|-------|-----|
| | PAT | CENT 1 | NO. | | | | | DATE | | | | | | | | D | ATE | |
| | | | | | | | | | | | | | | | | | | |
| PΙ | WO | 2004 | 0625 | 75 | | A2 | | 2004 | 0729 | , | WO 2 | 003-1 | JS41' | 720 | | 2 | 0031 | 222 |
| | WO | 2004 | 0625 | 75 | | A3 | | 2004 | 1209 | | | | | | | | | |
| | | W: | ΑE, | AG, | AL, | AM, | ΑT, | AU, | ΑZ, | BA, | BB, | BG, | BR, | BY, | ΒZ, | CA, | CH, | CN, |
| | | | CO, | CR, | CU, | CZ, | DE, | DK, | DM, | DΖ, | EC, | EE, | EG, | ES, | FI, | GB, | GD, | GE, |
| | | | GH, | GM, | HR, | HU, | ID, | ΙL, | IN, | IS, | JΡ, | ΚE, | KG, | ΚP, | KR, | ΚZ, | LC, | LK, |
| | | | LR, | LS, | LT, | LU, | LV, | MA, | MD, | MG, | MK, | MN, | MW, | MX, | MZ, | NI, | NO, | NZ, |
| | | | OM, | PG, | PH, | PL, | PT, | RO, | RU, | SC, | SD, | SE, | SG, | SK, | SL, | SY, | TJ, | TM, |
| | | | TN, | TR, | TT, | TZ, | UA, | UG, | US, | UZ, | VC, | VN, | YU, | ZA, | ZM, | ZW | | |
| | | RW: | GH, | GM, | KE, | LS, | MW, | MZ, | SD, | SL, | SZ, | TZ, | UG, | ZM, | ZW, | AM, | ΑZ, | BY, |
| | | | KG, | ΚZ, | MD, | RU, | TJ, | TM, | ΑT, | BE, | BG, | CH, | CY, | CZ, | DE, | DK, | EE, | ES, |
| | | | FΙ, | FR, | GB, | GR, | HU, | ΙE, | ΙΤ, | LU, | MC, | NL, | PT, | RO, | SE, | SI, | SK, | TR, |
| | | | BF, | ВJ, | CF, | CG, | CI, | CM, | GA, | GN, | GQ, | GW, | ML, | MR, | NE, | SN, | TD, | TG |
| | CA | 2512 | 713 | _ | | AA | | 2004 | 0729 | | CA 2 | 003-2 | 2512 | 713 | | 2 | 00312 | 222 |
| | US | 2004 | 2098: | 23) | | A1 | | 2004 | 1021 | • | US 2 | 003- | 7431 | 11 | | 2 | 0031 | 222 |
| • | ĒΡ | 1590 | 357 | | | A2 | | 2005 | 1102 | | EP 2 | 003-8 | 31009 | 90 | | 2 | 0031 | 222 |
| | | R: | AT, | ΒE, | CH, | DE, | DK, | ES; | FR, | GB, | GR, | ΙT, | LI, | LU, | NL, | SE, | MC, | PT, |
| | | | ΙE, | SI, | LT, | LV, | FI, | RO, | MK, | CY, | AL, | TR, | BG, | CZ, | EE, | HU, | SK | |
| PRAI | US | 2003 | -438 | 475P | | P | | 2003 | 0107 | | | | | | | | | |
| | WO | 2003 | -US4 | 1720 | | W | | 2003 | 1222 | | | | | | | | | |
| OS | MAR | PAT | 141: | 1340 | 50 | | | | | | | | | | | | | |
| GI | | | | | | | | | | | | | | | | | | |

Ι

AB We describe the use of bicyclic carbohydrates for the treatment of hepatitis C virus infections. Different bicyclic carbohydrates were tested in vitro against DNA-viruses, retroviruses and Flaviviridae sp., an important family of human and animal RNA pathogens. Significant activity was observed against the bovine viral diarrhea virus (BVDV). As pestiviruses, such as BVDV, share many similarities with hepatitis C virus (HCV), the bicyclic carbohydrates in general and the preferred bicyclic carbohydrates more specifically are expected to be a treatment for hepatitis C viral infections. I (preparation given) had an EC50 <0.8 $\mu g/mL$ against BVDV.

IT 727415-90-9P 727415-91-0P 727415-92-1P RL: BSU (Biological study, unclassified); PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological

study); PREP (Preparation); USES (Uses)

(bicyclic carbohydrate compds. useful in treatment of infections caused by Flaviviridae sp., such as hepatitis C and bovine viral diarrhea viruses)

RN 727415-90-9 CAPLUS

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-1-deoxy-1-phenyl-5,7-O-[(R)-phenylmethylene]-3,4-di-O-propyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

RN 727415-91-0 CAPLUS

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-3,4-di-O-butyl-1-deoxy-1-phenyl-5,7-O-[(R)-phenylmethylene]- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

RN 727415-92-1 CAPLUS

CN D-Glucitol, 1,5-anhydro-2,3-bis-O-(1-methylethyl)-1-C-phenyl-4,6-O-[(R)-phenylmethylene]-, (1S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

IT 727415-94-3P 727415-96-5P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(bicyclic carbohydrate compds. useful in treatment of infections caused by Flaviviridae sp., such as hepatitis C and bovine viral diarrhea viruses)

RN 727415-94-3 CAPLUS

Absolute stereochemistry. Rotation (-).

RN 727415-96-5 CAPLUS

CN D-Glucitol, 1,5-anhydro-2,3-bis-O-(1-methylethenyl)-1-C-phenyl-4,6-O-[(R)-phenylmethylene]-, (1S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

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L4
     ANSWER 9 OF 15 CAPLUS COPYRIGHT 2006 ACS on STN
     2004:350722 CAPLUS
AN
DN
     141:332381
     Alkaline Degradation of Glycosides. Part/5. Synthesis of a Ring-Rigid
ΤI
     Disaccharide Model for Studies of Alkal/ne Chain Cleavage in Cellulose
ΑU
     Croll, David C.; Schroeder, Leland R.
     Paper Science and Engineering Faculty/SUNY College of Environmental
CS
     Science and Forestry, Syracuse, NY, $\frac{1}{3210}$, USA
SO
     Journal of Wood Chemistry and Technology (2004) 24(1), 27-38
     CODEN: JWCTDJ; ISSN: 0277-3813
PB
     Marcel Dekker, Inc.
DT
     Journal
LΑ
     English
     2,6-Anhydro-1,3-O-methylene-5-0/-(4,6-O-methylene-\beta-D-glucopyranosyl)-
     D-glycero-D-gulitol was synthesized as a non-reducing cellulose model
     compound for studying high temperature alkaline cleavage of the glucosidic
linkage.
     The pyranoid rings are limited in their ability to change conformation by
     the methylene moieties attached to each ring. Thus the compound more
     closely models the situat/on for internal glycosidic linkages of
     cellulose. A related model with the 2' hydroxyl group etherified,
     2,6-anhydro-1,3-0-methylene-5-0-(2-0-methyl-4,6-0-methylene-\beta-D-
     glucopyranosyl)-D-glycéro-D-gulitol was also synthesized.
     770727-51-0P 770727-53-2P
     RL: RCT (Reactant); $PN (Synthetic preparation); PREP (Preparation); RACT
```

RN 770727-51-0 CAPLUS

(Reactant or reagent/)

cleavage in cel/ulose)

(synthesis of a ring-rigid disaccharide model for studies of alkaline chain

Absolute stereochemistry. Rotation (+).

RN 770727-53-2 CAPLU\$

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-5-O-[2,3-bis-O-(phenylmethyl)-4,6-O-[(R)-phenylmethylene]-β-D-glucopyranosyl]-4,7-bis-O-(phenylmethyl)-1,3-O-[(R)-phenylmethylene]- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (+).

RE.CNT 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 10 OF 15 CAPLUS COPYRIGHT 2006 AC\$ on STN

AN 2003:738805 CAPLUS

DN 139:381689

TI Synthesis of Both Possible Isomers of the Northwest Quadrant of Altromycin

AU Pasetto, Paolo; Franck, Richard W.

CS Department of Chemistry, Hunter College The City University of New York, New York, NY, 10021, USA

SO Journal of Organic Chemistry (2003), 68(21), 8042-8060 CODEN: JOCEAH; ISSN: 0022-3263

PB American Chemical Society

DT Journal

LA English

OS CASREACT 139:381689

AB The synthesis of the northwest quadrant of Altromycin B is described. The preparation of the two epimers at the quaternary carbon of the 6-deoxy-C-altrose moiety in the northwest quadrant is accomplished starting from D-glucose. A key step of our synthetic sequence is the formation of the C-glycoside linkage via the Ramberg-Backlund reaction. Two different routes are explored, which differ mainly on the timing of the conversion of glucose to altrose, either before or after the preparation of the C-glycoside. The conformation behavior of variously substituted C-altropyranoside rings is also discussed.

IT 623148-57-2P 623148-59-4P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(synthesis and conformation of both possible isomers of northwest quadrant of altromycin B from D-glacose via Ramberg-Backlund reaction)

RN 623148-57-2 CAPLUS

CN D-glycero-L-gulo-Heptitol, 2,6-anhydro-4,5-bis-O-[(4-methoxyphenyl)methyl]-7-C-phenyl-1,3-O-[(R)-phenylmethylene]-, (7R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 623148-59-4 CAPLUS

CN D-glycero-L-gulo-Heptitol, 2,6-anhydro-4,5-bis-O-[(4-methoxyphenyl)methyl]-7-C-phenyl-1,3-O-[(R)-phenylmethylene]-7-O-[[2-(trimethylsilyl)ethoxy]methyl]-, (7R)- (9CI) (CA INDEX NAME)

IT 623148-58-3P

RN

RL: SPN (Synthetic preparation); PREP (Preparation)
(synthesis and conformation of both possible isomers of northwest
quadrant of altromycin B from D-glucose via Ramberg-Backlund reaction)
623148-58-3 CAPLUS

CN D-glycero-D-gylo-Heptitol, 2,6-anhydro-3,4-bis-O-[(4-methoxyphenyl)methyl]-1-C-phenyl-5,7-O-[(R)-phenylmethylene]-, (1S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RE.CNT 78 THERE ARE 78 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 11 OF 15 CAPLUS COPYRIGHT 200% ACS on STN

AN 2003:595156 CAPLUS

DN 140:104492

TI Analogues of the mycobacterial arabinogalactan linkage disaccharide as cell wall biosynthesis inhibitors

AU Wen, Xianghui; Crick, Dean C.; Brennan, Patrick J.; Hultin, Philip G.

CS Department of Chemistry, University of Manitoba, Winnipeg, MB, R3T 2N2,

SO Bioorganic & Medicinal Chemistry (2003), 11(17), 3579-3587 CODEN: BMECEP; ISSN: 0968-0896 PB Elsevier Science Ltd.

DT Journal

LA English

AB The mycobacterial arabinogalactan linkage disaccharide [α-L-Rha-(1→3)-α-D-GlcNAc] provides a basis for the design of new antitubercular drugs, since it supports a key skeletal structure in the bacterial cell wall. A series of analogs of the linker was synthesized by coupling appropriate thiorhamnosyl donors modified at their 4-positions, with an N-acetyl glucosamine acceptor. In a cell-free enzyme inhibition assay, three analogs inhibited the activity of the galactosyltransferase that adds a Galf residue to the linkage disaccharide. Although the compds. Were modest inhibitors, these data confirm the viability of this approach to anti-mycobacterial agents. It is especially significant that the three effective compds. are modified at the site of the acceptor atom in the natural substrate.

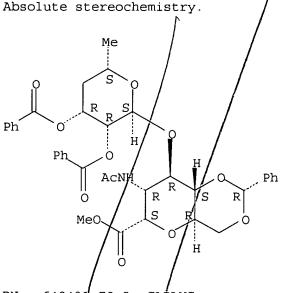
IT 648431-78-1P 648431-79-2P 648431-84-9P

RL: PRP (Properties); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(analogs of the mycobacterial arabinogalactan linkage disaccharide as cell wall biosynthesis inhibitors)

RN 648431-78-1 CAPLUS

CN D-glycero-D-ido-Heptonic acid, 3-(acetylamino)-2,6-anhydro-3-deoxy-4-0-(2,3-di-0-benzoyl-4,6-dideoxy-α-L-lyxo-hexopyranosyl)-5,7-0-[(R)-phenylmethylene]-, methyl ester (9CI) (CA INDEX NAME)



RN 648431-79-2 CAPLUS

CN

D-glycero-D-ido-Heptonic acid, 3-(acetylamino)-2,6-anhydro-3-deoxy-4-0-(2,3-di-0-benzoyl-6-deoxy-4-0-methyl- α -L-mannopyranosyl)-5,7-0-[(R)-phenylmethylene]-, methyl ester (9CI) (CA INDEX NAME)

RN 648431-84-9 CAPLUS

CN D-glycero-L-gulo-Heptitol, 5-(acetylamino)-2,6-anhydro-5,7-dideoxy-4-O-(2,3-di-O-benzoyl-6-deoxy-4-O-methyl-α-L-mannopyranosyl)-7-(diethoxyphosphinyl)-1,3-O-[(R)-phenylmethylene]-(9CI) (CA INDEX NAME)

RE.CNT 39 THERE ARE 39 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L4 ANSWER 12 OF 15 CAPLUS COPYRIGHT 2006 ACS on STN
- AN 2003:384377 CAPLUS
- DN 139:230897
- TI Synthesis of thio-C-glycosides from 2'-carbonylalkyl C-glycosides by a tandem β -elimination and intramolecular/hetero-Michael addition
- AU Zou, Wei; Lacroix, Edith; Wang, Zerong; Wu, Shih-Hsiung
- CS Institute for Biological Sciences, National Research Council of Canada, Ottawa, ON, K1A 0R6, Can.
- SO Tetrahedron Letters (2003), 44(24), 4431-4433 CODEN: TELEAY; ISSN: 0040-4039
- PB Elsevier Science Ltd.
- DT Journal
- LA English
- OS CASREACT 139:230897
- AB 2'-Carbonyl 5-S-acetyl-C-glycofuranosides and 2'-carbonyl

```
4-S-acetyl-C-glycopyranosides were/converted in good yields to
     5-thio-C-glycopyranosides and 4-thio-C-glycofuranosides under base
     treatment. The transformation résulted from \beta-elimination on
     2'-carbonyl C-glycoside to form/\alpha, \beta-conjugated aldehyde (or
     ketone) and subsequent intramo/. hetero-Michael addition by the thiol group.
ΙT
     592532-99-5P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation of thio-C-glycosides from 2'-carbonylalkyl C-glycosides via
        tandem \beta-elimination and intramol. hetero-Michael addition as the key
        steps)
     592532-99-5 CAPLUS
RN
CN
     D-glycero-L-galacto-Non-8/enitol, 2,6-anhydro-7,8,9-trideoxy-4,5-bis-O-
     (phenylmethyl)-1,3-O-[(S)/-phenylmethylene]- (9CI) (CA INDEX NAME)
Absolute stereochemistry.
           R
                   Ph
              THERE ARE/14 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 14
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
     ANSWER 13 OF 15 CAPLUS COPYRIGHT 2006 ACS on STN
L4
AN
     2003:319645 CAPLUS
DN
     138:321501
ΤI
     Preparation and use of glycoside-based ring structures with antimicrobial
     and cytostatic activity
IN
     Sas, Benedikt; Van Der Eycken, Johan; Van Hemel, Johan; Blom, Petra;
     Vandenkerckhove, Jan; Ruttens, Bart
PΑ
     Kemin Pharma Europe, USA
                                        Pork
     PCT Int. Appl., 99 pp.
SO
     CODEN: PIXXD2
DT
     Patent
LΑ
     English
FAN.CNT 2
     PATENT NO.
                         KIND
                                 DATE
                                             APPLICATION NO
                                                                     DATE
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                                 _____
ΡI
     WO 2003032905
                          A2
                                 20030424
                                             WO 2002-US 22817
                                                                     20021015
     WO 2003032905
                          Α3
                                 20040129
            AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, Ex, ES, FI, GB, GD, GE, GH,
             GM, HR, HU, ID, IL, IN, IS, ar{\mathsf{JP}}, KE, ar{\mathsf{K}}G, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN/MW, MX, MZ, NO, NZ, OM, PH,
             PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,
             UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
             FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF,
             CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
     US 2003158243
                          Α1
                                 20030821
                                             US 2001-977478
                                                                     20011015
     CA 2463084
                          AΑ
                                 20030424
                                             CA 2002-2463084
                                                                     20021015
     EP 1446391
                          A2
                                 20040818
                                             EP 2002-770578
                                                                     20021015
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AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,

IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK

| | JP 2005509615 | T2 | 20050414 | JP 2003-535711 | 20021015 |
|------|-------------------|----|----------|----------------|----------|
| | US 2004224904 | A1 | 20041111 | US 2004-861768 | 20040604 |
| PRAI | US 2001-977478 | Α | 20011015 | | |
| | WO 2002-US32817 | W | 20021015 | | |
| os | MARPAT 138:321501 | | | | |
| GI | | | | | |

Novel glycoside-based compds. I, wherein R1 is H, SPh, Ph, PhS, All, Bn; R2 is H, Et, All, Me, Bn; R3 is H, Et, Me, All, Bn; R4 and R5 form a ring and are -carbamate-C6-alkyl-ether-C4-alkenyl-ether-, -ester-C6-alkyl-ester-, -ether-C8-alkenyl-ether-, -ester-C6-alkenyl-amide-, -ether-C7-alkenyl-amide-, -ester-C10-alkenyl-ester-, -ester-C18-alkenyl-ester-, -OCH(Ph)CH2O-, with an attached ring system that have antimicrobial or cytostatic activity. The compds. are administered to humans and animals for the treatment or amelioration of bacterial, fungal, viral, or protozoal infections or tumors. Thus, glycoside II was prepared and tested in humans for its antimicrobial and cytostatic activities.

IT 511274-41-2P 511274-42-3P

RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)

(preparation and use of glycoside-based ring structures with antimicrobial and cytostatic activity)

RN 511274-41-2 CAPLUS

CN D-Glucitol, 1,5-anhydro-2,3-di-O-methyl-1-C-phenyl-4,6-O-[(R)-phenylmethylene]-, (1S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

RN 511274-42-3 CAPLUS

CN D-glycero-D-gulo-Heptitol, 2,6-anhydro-1-deoxy-3,4-di-O-methyl-1-phenyl-5,7-O-[(R)-phenylmethylene]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

IT 511274-46-7P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation and use of glycoside-based ring structures with antimicrobial and cytostatic activity)

RN 511274-46-7 CAPLUS

CN D-glycero-D-gulo-Non-1-enitol, 4,8-anhydro-1,2,3-trideoxy-5,6-di-O-methyl-7,9-O-[(R)-phenylmethylene]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

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L4 ANSWER 14 OF 15 CAPLUS COPYRIGHT 2006 ACS on STN
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AN 2002:859447 CAPLUS

DN 138:89617

TI Convergent synthesis of the BCDE-ring part of ciguatoxin

AU Fujiwara, Kenshu; Koyama, Yasuhito; Kawai, Kentaro; Tanaka, Hideki; Murai, Akio

CS Division of Chemistry, Graduate School of Science, Hokkaido University, Sapporo, 060-0810/ Japan

SO Synlett (2002), (11), 1835-1838 CODEN: SYNLES; ISSN: 0936-5214

PB Georg Thieme Verlag

DT Journal

LA English

OS CASREACT 138:8/9617

GI

AB The BCDE-ring part of ciguatoxin, I, has been convergently synthesized by a sequence of reactions involving coupling of the E-ring part, II, having a Z-enal group with the B-ring part, III, as an acyl anion equivalent, selective removal of a p-bromobenzyl protective group using Miyaura's conditions, and reductive cyclization.

IT 482646-97-9

RL: RCT (Reactant); RACT (Reactant or reagent)

(stered selective preparation of BCDE ring system of ciquatoxin)

RN 482646-97**f**9 CAPLUS

CN D-glycero-D-gulo-Octitol, 3,7-anhydro-2-deoxy-6,8-0-[(R)-(4-

methoxyphenyl)methylene]-4,5-bis-O-(phenylmethyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

IT 482646-98-0P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(stereoselective preparation of BCDE ring system of ciguatoxin)

RN 482646-98-0 CAPLUS

CN D-glycero-D-gulo-Octitol, 3,7-anhydro-2-deoxy-6,8-O-[(R)-(4-methoxyphenyl)methylene]-1,4,5-tris-O-(phenylmethyl)- (9CI) (CA INDEX NAME)

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OMe
             Ph
                  S
RE.CNT 63
               THERE ARE 63 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
     ANSWER 15 OF 15 CAPLUS COPYRIGHT 2006 ACS on STN
L4
AN
     1946:33217 CAPLUS
DN
     40:33217
OREF 40:6418e-i,6419a
     Carbohydrate C-nitro alcohols: 1-\text{nitro}/1-\text{desoxy-D-}\alpha-\text{glucoheptitol}
     and a 2,6-anhydro-1-nitro-1-desoxyheptitol
ΑU
     Sowden, John C.; Fischer, Hermann O.
CS
     Univ. of Toronto, Can.
SO
     Journal of the American Chemical Society (1946), 68, 1511-13
     CODEN: JACSAT; ISSN: 0002-7863
DT
     Journal
LΑ
     Unavailable
     cf. C.A. 40, 61.9. 4,6-Benzyliden glucose (I) (100 g.) in 1500 cc. absolute
AΒ
     MeOH and 250 cc. MeNO2, treated w∳th 17.3 g. Na in 1 l. MeOH, allowed to
     stand at room temperature for 8 h\!\!\!/ xs., acidified with 47 cc. AcOH, evaporated
to a
     thin sirup, 200 cc. H2O added, and the solution again concentrated, gives 21%
of
     the 5,7-benzylidene derivative /(II), m. 160-2°, [\alpha]D28
     -42.5° (H2O, c 1.1), of 1-nitro-1-desoxy-D-glucoheptitol (III), m.
     138-9°, [\alpha] D23 3.1° (H2O, c 6./4); HI results in 77%
     yield on hydrolysis of 5 g. of II with 50 cc. 0.1 N H2SO4 at 70-5°
     for 45 min. and at 100° for 1/5 min. Both II and III vigorously
     reduce hot Fehling solution / The structure of III follows from the conversion
     of its Na salt by dilute H2SO4 into D-\alpha-gLucoheptose (isolated as the
     benzylphenylhydrazone). III (2 g.) in 40 cc. H2O, reduced over 1 q. Raney
     Ni for 55 min., gives 1-amiho-1-desoxy-D-\alpha-glucoheptitol, isolated
     as the p-toluenesulfonate, \int_{0}^{\infty} 140-50^{\circ}, [\alpha] D22 -4.2°
     (H2O, c 4),84% yield. I (\rlap/26g.),250 cc. MeNO2, 3160 cc. MeOH, and 21.7 g.
     Na, allowed to stand at r\phiom temperature for 24 hrs. and acidified with AcOH,
     give a sirupy product containing 5% of the 5,7-benzylidene derivative (IV), m.
     211-12°, [α]D31 -35.4° (MeOH, c 1.3), of
     2,6-anhydro-1-nitro-1-de\phioxyheptitol (V), m. 177-7.5°, [\alpha] D22
     8.2° (H2O, c 3.7). If the formation of IV is carried out at
     6-8° for 18 days, there presults 3% of IV and 19% of II. IV and V
     are indifferent to hot Fehling solution and OsO4. With 0.1 N Pb(OAc)4 in 98%
     AcOH, IV consumes 0.96 mol. of oxidant in 68 hrs. and 1.22 mols. in 115
     hrs. IV with Ac2O and AcONa (5 min. at 100°) yields 69% of the
     diacetate, m. 192-3°, [\alpha]D23 -60° (CHCl3, c 2). V
     with Ac20 containing a/trace of H2SO4 (2 days at room temperature) yields a
     tetraacetate, m. 144-5\beta, [\alpha] D23 4.2^{\circ} (CHCl3, c 4.2).
     With Na metaperiodate / V consumes 2 equivs. of oxidant and yields 1 equivalent
     of HCO2H but no HCHO. / The 2,6-position is therefore assigned to the
     anhydro ring. The 2 acetates are not affected by solid NaHCO3 in boiling
     C6H6, indicating the absence of an Ac group on the C atom adjacent to the
     CH2NO2 group.
IT
     867219-64-5, Heptitol, 5,7-benzylidene-2,6-anhydro-1-nitro-1-
     desoxy-, diacetate
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(preparation of)

RN 867219-64-5 CAPI/US Heptitol, 5,7-be/zylidene-2,6-anhydro-1-nitro-1-desoxy-, diacetate (4CI) CN (CA INDEX NAME) CH2 - NO2 OAd ÖAc => file toxicenter 'TOXICENTER' IS NOT A VALID FILE NAME

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